

safety and quality standards. The Australian Commission on Safety and Quality in Health Care has developed resources that assist health services implement these important initiatives.

The introduction of these requirements have been 'transformative' in that it provides the necessary stimulus for hospital executives to re-prioritise and commit resources to the prevention of healthcare associated infections as well as accept responsibility for governance.

Chen AWJ, Khumra S, Eaton V, Kong D Snapshot of antimicrobial stewardship in Australian hospitals Journal of Pharmacy Practice and Research Volume 41, No. 1, 2011

#### SP 6-2

##### SURVEILLANCE SYSTEM OF ANTIMICROBIAL RESISTANCE AND HEALTHCARE-ASSOCIATED INFECTIONS IN JAPAN

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Antimicrobial resistance (AMR) is of global concern, and strengthening surveillance is critical. Japan has two major surveillance for AMR. The first one is National Epidemiological Surveillance of Infectious Disease (NESID), which is a national comprehensive surveillance system under infection control act. About 110 infectious diseases including seven AMR bacterial infections are designated as reportable disease. The second one is Japan Nosocomial Infections Surveillance (JANIS), which has been established in 2000 as a voluntary-based national surveillance system targeting healthcare-associated infections and antimicrobial resistant bacteria. It is organized by Ministry of Health Labour and Welfare, and JANIS management office located at National Institute of Infectious Diseases. More than 1600 hospitals are participating to JANIS on voluntary basis as of January 1<sup>st</sup>, 2015.

JANIS consist of five divisions and Clinical Laboratory division provides information on prevalence of AMR bacteria among clinical isolates. JANIS collects data from hospital laboratories that have an automated system for bacterial identification and drug susceptibility testing, and also data from commercial laboratories to which participating hospitals are contracted. Following conversion of those laboratory data into JANIS data format, the data is collected to a centralized JANIS database server for analysis. Collected data would be processed by JANIS tabulation program, which was developed by JANIS research team, to detect data error, remove duplicate samples and differentiate drug susceptibility pattern by an original algorithm. JANIS has been working for years to standardize JANIS data format and establish tabulation program. The outputs from this JANIS system include aggregate AMR data by hospital to allow for inter-hospital comparison as well as national data for AMR trends. From 2014, JANIS starts to recruit hospitals outside Japan to participate this surveillance system.

In the presentation, a summary of JANIS annual report 2013 which is based on data of 4.6 million samples and 3.6 million isolates submitted from 745 hospitals across Japan would be provided.

#### SP 6-3

##### IMPLEMENTATION OF BUNDLE CARE TO DECREASE HEALTHCARE ASSOCIATED INFECTIONS IN TAIWAN

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Health care-associated infections (HAI) are associated with significant morbidity at hospitals. We implemented active surveillance, hand hygiene promotion, isolation cohort of resistant microorganism and bundle care since 2010. The bundle care program includes prevention care of center-line associated bloodstream infection, urinary catheter associated urinary tract infection, ventilator-associated pneumonia, and surgical wound infection. The infection density decreased from 5.1 per 1000 patient-days in 2011, 4.9 in 2012, 4.2 in 2013, and 3.7 in 2014, respectively. Infection density of methicillin-resistant *Staphylococcus aureus* had decreased from 0.18 per 1000 patient-days in 2011 to 0.11 in 2014. Infection density of carbapenem-resistant *Acinetobacter baumannii* had decreased from 0.15 per 1000 patient-days (‰) in

2011, 0.18 in 2012, 0.09 in 2013 and 0.07 in 2014 ( $P < 0.001$ ). A infection control environment reduce the morbidity, mortality and medical costs. Implementation of bundle care has shown successful to decrease HAI in Taiwan.

#### SYMPOSIUM 7 (SP 7)

##### CLOSTRIDIUM DIFFICILE INFECTIONS (CDI)

#### SP 7-1

##### CLOSTRIDIUM DIFFICILE INFECTION IN WESTERN COUNTRIES: UNDER DIAGNOSES, EPIDEMIOLOGY & EVOLVING THERAPIES

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*C. difficile* associated infection (CDI) effects approximately 500–700,000 Americans annually and is associated with significant and recently increasing morbidity and mortality with 7,285 deaths reported in 2009. CDI also has a substantial economic impact. The cost of treating one CDI patient in hospital is approximately \$8,000. Annual expenditures in the USA to manage CDI is \$3.2 billion/year, not including measures taken to prevent the spread of *C. difficile* spores. Because of lack of suspicion and/or laboratory methodological issues it seems likely that CDI is underdiagnosed worldwide. It is estimated that 40,000 cases per year are missed in European countries. Of 336,600 hospitalizations in the US, 1% of all stays develop CDI. Rates vary for HO-HCFA CDI from rates 2.8–9.3 per 10,000 patient days, to 1.3–2.7 CO-HCFA and 20–30 per 100,000 population for CA-CDI. In the USA stool culture for *C. difficile* is not performed and therefore epidemiological surveys rely on data from a few centers. Between 2011–2013, 29 ribotypes types were identified with 027 most frequent and accounting for 28.1% although rates varied by geographic region. In a survey of 20 European countries, 027 was the most prevalent ribotype (~18%) but varied by country for 43% in Germany to 12% in Romania.

The therapy for CDI has been stagnant for approximately 30 years with the only alternatives being metronidazole or oral vancomycin. Because relapse rates range between 20–30%, especially in the elderly and medically vulnerable, there has been a search for both new and more effective therapeutic agents as well as novel preventative strategies. A recent study (CID 2014) notes the inferiority of both metronidazole and vancomycin compared to fidaxomicin in mild, moderate and severe CDI with statistically superior sustained cure especially in various patient groups at greater risk of relapse such as patients requiring concomitant antibiotics and those with renal failure. Several new agents surotomycin, (CB-183315), SMT19969, cadazolid and others are currently in development for CDI therapy. Other promising alternative therapeutic approaches include monoclonal antibodies, fecal biotherapy, oral spore ingestion, vaccination, and the use of probiotics for primary prevention.

#### SP 7-2

##### HYPERVIRULENT CLOSTRIDIUM DIFFICILE IN ASIA

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While *Clostridium difficile* infection (CDI) has come to prominence as major epidemics have occurred in North America and Europe over the last 15 years, awareness and surveillance of CDI in Asia have remained poor. Limited studies performed throughout Asia indicate that CDI is also a significant nosocomial pathogen in this region, but the true prevalence of CDI remains unknown. A lack of regulated antibiotic use in many Asian countries suggests that the prevalence of CDI may be comparatively high. Molecular studies indicate that ribotypes 027 and 078, which have caused significant outbreaks in other regions of the world, are rare in human CDI in Asia, however, there have been no published investigations of production or companion animals in the region. Variant toxin A-negative/toxin B-positive strains of ribotype 017 caused apparent epidemics across several Asian countries in the mid-2000s but now appear to be waning. Ribotype smz/018 has caused widespread disease across Japan over the last decade and more recently emerged in Korea. Both ribotype 17 and 18 have caused major outbreaks outside Asia highlighting the potential for transfer to different countries. Better surveillance for CDI in Asia is essential, and urgently required.